AMENDMENTS TO THE CLAIMS

The following listing of claims replaces all prior versions, and listings, of claims in this application.

Listing of Claims:

1. (Previously Presented) A compound having the formula:

or a salt thereof.

- 2. (Previously Presented) A composition comprising:
 - (A) an active agent; and
 - (B) the compound of claim 1.
- 3. (Original) The composition of claim 2, wherein the active agent is selected from the group consisting of a biologically active agent, a chemically active agent, and a combination thereof.
- 4. (Previously Presented) The composition of claim 3, wherein the biologically active agent comprises at least one protein, polypeptide, peptide, hormone, polysaccharide, mucopolysaccharide, carbohydrate, or lipid.
- 5. (Original) The composition of claim 3, wherein the biologically active agent is selected from the group consisting of: growth hormones, human growth hormones recombinant human growth hormones (rhGH), bovine growth hormones, porcine growth hormones, growth hormone releasing hormones, growth hormone releasing factor, interferons, α -interferon, β -

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interferon, γ-interferon, interleukin-1, interleukin-2, insulin, porcine insulin, bovine insulin, human insulin, human recombinant insulin, insulin-like growth factor (IGF), IGF-1, heparin, unfractionated heparin, heparinoids, dermatans, chondroitins, low molecular weight heparin, very low molecular weight heparin, ultra low molecular weight heparin, calcitonin, salmon calcitonin, eel calcitonin, human calcitonin; erythropoietin (EPO), atrial naturetic factor, antigens, monoclonal antibodies, somatostatin, protease inhibitors, adrenocorticotropin, gonadotropin releasing hormone, oxytocin, leutinizing-hormone-releasing-hormone, follicle stimulating hormone, glucocerebrosidase, thrombopoeitin, filgrastim. postaglandins, cyclosporin, vasopressin, cromolym sodium, sodium chromoglycate, disodium chromoglycate, vancomycin, desferrioxamine (DFO), parathyroid hormone (PTH), fragments of PTH, antimicrobials, anti-fungal agents, vitamins; analogs, fragments, mimetics and polyethylene glycol (PEG)-modified derivatives of these compounds; and any combination thereof.

- 6. (Original) The composition of claim 3, wherein the biologically active agent comprises insulin, heparin, calcitonin, parathyroid hormone, erythropoietin, growth hormones or combinations thereof.
- 7. (Previously Presented) The composition of claim 3, wherein the biologically active agent comprises a recombinant human growth hormone.
- 8. (Original) The composition of claim 3, wherein the biologically active agent comprises parathyroid hormone.
- 9. (Original) The composition of claim 3, wherein the biologically active agent comprises insulin.
- 10. (Original) The composition of claim 3, wherein the biologically active agent comprises heparin.
- 11. (Original) The composition of claim 3, wherein the biologically active agent comprises calcitonin.

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12. (Original) The composition of claim 3, wherein the biologically active agent comprises interferon.

13-14 (Canceled)

- 15. (Previously Presented) A dosage unit form comprising:
 - (A) the composition of claim 2; and
 - (B) (a) an excipient,
 - (b) a diluent,
 - (c) a disintegrant,
 - (d) a lubricant,
 - (e) a plasticizer,
 - (f) a colorant,
 - (g) a dosing vehicle, or
 - (h) any combination thereof.
- 16. (Original) The dosage unit form of claim 15, wherein the active agent is selected from the group consisting of a biologically active agent, a chemically active agent, and a combination thereof.
- 17. (Original) The dosage unit form of claim 16, wherein the biologically active agent comprises at least one protein, polypeptide, peptide, hormone, polysaccharide, mucopolysaccharide, carbohydrate, or lipid.
- 18. (Original) The dosage unit form of claim 16, wherein the biologically active agent is selected from the group consisting of: growth hormones, human growth hormones (hGH), recombinant human growth hormones (rhGH), bovine growth hormones, porcine growth hormones, growth hormone releasing hormones, growth hormone releasing factor, interferons, α-interferon, β-interferon, γ-interferon, interleukin-1, interleukin-2, insulin, porcine insulin, bovine insulin, human insulin, human recombinant insulin, insulin-like growth factor, insulin-like growth factor-1, heparin, unfractionated heparin, heparinoids,

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dermatans, chondroitins, low molecular weight heparin, very low molecular weight heparin, ultra low molecular weight heparin, calcitonin, salmon calcitonin, eel calcitonin, human calcitonin; erythropoietin, atrial naturetic factor, antigens, monoclonal antibodies, somatostatin, protease inhibitors, adrenocorticotropin, gonadotropin releasing hormone, oxytocin, leutinizing-hormone-releasing-hormone, follicle stimulating hormone, glucocerebrosidase, thrombopoeitin, filgrastim. postaglandins, cyclosporin, vasopressin, cromolym sodium, sodium chromoglycate, disodium chromoglycate, vancomycin, desferrioxamine, parathyroid hormone, fragments of PTH, antimicrobials, anti-fungal agents, vitamins; analogs, fragments, mimetics and polyethylene glycol-modified derivatives of these compounds; and any combination thereof.

- 19. (Original) The dosage unit form of claim 16, wherein the biologically active agent comprises insulin, heparin, calcitonin, parathyroid hormone, erythropoietin, human growth hormones or combinations thereof.
- 20. (Original) The dosage unit form of claim 15, wherein the active agent comprises recombinant human growth hormone.
- 21. (Original) The dosage unit form of claim 15, wherein the active agent comprises parathyroid hormone.
- 22. (Original) The dosage unit form of claim 15, wherein the active agent comprises insulin.
- 23. (Original) The dosage unit form of claim 15, wherein the active agent comprises heparin.
- 24. (Original) The dosage unit form of claim 15, wherein the active agent comprises calcitonin.
- 25. (Original) The dosage unit form of claim 15, wherein the active agent comprises interferon.

- 26. (Original) The dosage unit form of claim 15, wherein the dosage unit form comprises a dosing vehicle comprising a tablet, a capsule, a powder, or a liquid.
- 27. (Previously Presented) The dosage unit form of claim 15, wherein the dosing vehicle is <u>a</u> liquid selected from the group consisting or water, 1,2-propane diol, ethanol, and any combination thereof.
- 28. (Original) A method for administering a biologically-active agent to an animal in need of the agent, the method comprising administering orally to the animal the composition of claim 3.
 - 29. (Original) A method for preparing a composition comprising mixing:
 - (A) at least one active agent;
 - (B) the compound of claim 1; and
 - (C) optionally, a dosing vehicle.
- 30. (Previously Presented) The composition of claim 4, wherein the biologically active agent comprises a peptide.
- 31. (Previously Presented) The dosage unit form of claim 17, wherein the biologically active agent comprises a peptide.